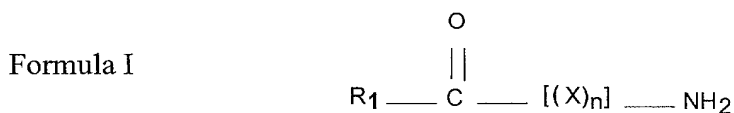


We claim:

1. An antimicrobial composition comprising a plurality of hexapeptides
wherein for each hexapeptide, the amino acid in the first position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, lysine, methionine, serine, threonine and tryptophan;
the amino acid in the second position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, histidine, cysteine, threonine, tyrosine, and tryptophan;
the amino acids in positions three through six, based on numbered amino acids from N-terminus to C-terminus, are any amino acid; and
wherein the first two amino acids of said hexapeptides are other than arginine-arginine, tryptophan-tryptophan, tryptophan-cysteine, tryptophan-lysine, arginine-tryptophan, or threonine-arginine.
2. The antimicrobial composition of claim 1 wherein the amino acids in the first and second positions of said peptides, based on numbered amino acids from N-terminus to C-terminus, are selected from the group consisting of Arg-Tyr, Arg-Cys, Ser-Thr, Met-Trp, Lys-Trp, Thr-Trp, Trp-Arg, Trp-His, and Trp-Tyr.
3. The antimicrobial composition of claim 1 wherein said peptides are incorporated into a polymer.
4. The antimicrobial composition of claim 3 wherein said polymer is selected from the group consisting of a polysaccharide, a glycol polymer, a polyester, a polyurethane, a polyacrylate, a polyacrylonitrile, a polyamide, a polyolefin, a polystyrene, a vinyl polymer, a polypropylene, silk, a biopolymer, and mixtures thereof.
5. An antimicrobial composition comprising a plurality of peptides, wherein said peptides

each are represented by Formula I:



wherein:

X represents any amino acid except glutamate or aspartate;

n = 6;

R₁ is C₁-C₂₀ alkyl; C₃-C₆ cycloalkyl; C₄-C₂₀ alkenyl; C₄-C₂₀ alkynyl; C₁-C₂₀ haloalkyl; C₃-C₂₀ haloalkenyl; C₃-C₂₀ haloalkynyl; C₂-C₂₀ alkoxyalkyl; C₂-C₂₀ alkylthioalkyl; C₂-C₂₀ alkylsulfinylalkyl; C₂-C₂₀ alkylsulfonylalkyl; C₅-C₂₀ cycloalkylalkyl; C₄-C₂₀ alkenyloxyalkyl; C₄-C₂₀ alkynyloxyalkyl; C₄-C₂₀ (cycloalkyl) oxyalkyl; C₄-C₂₀ alkenylthioalkyl; C₄-C₂₀ alkynylthioalkyl; C₆-C₂₀ (cycloalkyl) thioalkyl; C₂-C₂₀ haloalkoxyalkyl; C₄-C₂₀ haloalkenyloxyalkyl; C₄-C₂₀ haloalkynyloxyalkyl; C₄-C₂₀ alkoxyalkenyl; C₄-C₂₀ alkoxyalkynyl; C₄-C₂₀ alkylthioalkenyl; C₄-C₂₀ alkylthioalkynyl; C₄-C₂₀ trialkylsilylalkyl; C₁-C₂₀ alkyl substituted with NR₃R₄, nitro, cyano, or phenyl optionally substituted with R₅, R₆, and R₇; C₁-C₂₀ alkoxy; C₁-C₂₀ haloalkoxy; C₁-C₂₀ alkylthio; C₁-C₂₀ haloalkylthio; NR₃R₄; or phenyl, benzyl, pyridyl, furanyl, thienyl, naphthyl, pyrimidinyl, benzofuranyl, benzothienyl, or quinolinyl each optionally substituted with R₅, R₆ or R₇;

R₃ is independently hydrogen; C₁-C₄ alkyl; or phenyl optionally substituted with at least one R₈;

R₄ is independently hydrogen; C₁-C₈ alkyl; or phenyl optionally substituted with at least one R₈;

R₅ is independently C₁-C₆ alkyl; C₁-C₆ alkoxy; C₁-C₆ haloalkyl; halogen; C₂-C₈ alkynyl; C₁-C₆ thioalkyl; phenyl or phenoxy each optionally substituted with at least one R₈; cyano; nitro; C₁-C₆ haloalkoxy; C₁-C₆ haloalkylthio; C₂-C₆ alkenyl; C₂-C₆ haloalkenyl; acetyl; CO₂CH₃; or N(C₁-C₂ alkyl)₂;

R₆ is independently methyl; ethyl; methoxy; methylthio; halogen; or trifluoromethyl;

R₇ is independently halogen; and

R₈ is independently halogen; C₁-C₄ alkyl; C₁-C₄ alkoxy; C₁-C₄ haloalkyl; nitro; or cyano;

wherein:

the amino acid in the first position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, lysine, methionine, serine, threonine and tryptophan;

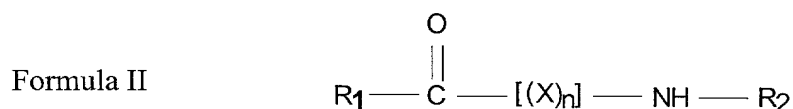
the amino acid in the second position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, histidine, cysteine, threonine, tyrosine, and tryptophan; and

the amino acids in positions three through six, based on numbered amino acids from N-terminus to C-terminus, are any amino acid;

wherein the first two amino acids of said hexapeptides are other than arginine arginine, tryptophan-tryptophan, tryptophan-cysteine, tryptophan-lysine, arginine-tryptophan, or threonine-arginine.

6. The antimicrobial composition of claim 5 wherein the amino acids in the first and second positions of said peptides, based on numbered amino acids from N-terminus to C-terminus, are selected from the group consisting of Arg-Tyr, Arg-Cys, Ser-Thr, Met-Trp, Lys-Trp, Thr-Trp, Trp-Arg, Trp-His, and Trp-Tyr.
7. The antimicrobial composition of claim 5 wherein said peptides are incorporated into a polymer.
8. The antimicrobial composition of claim 7 wherein said polymer is selected from the group consisting of a polysaccharide, a glycol polymer, a polyester, a polyurethane, a polyacrylate, a polyacrylonitrile, a polyamide, a polyolefin, a polystyrene, a vinyl polymer, a polypropylene, silk, a biopolymer, and mixtures thereof.
9. An antimicrobial composition comprising a plurality of peptides, wherein said peptides

each are represented by Formula II:



wherein:

X represents any amino acid except glutamate or aspartate;

n = 6;

R₁ is C₁-C₂₀ alkyl; C₃-C₆ cycloalkyl; C₄-C₂₀ alkenyl; C₄-C₂₀ alkynyl; C₁-C₂₀ haloalkyl; C₃-C₂₀ haloalkenyl; C₃-C₂₀ haloalkynyl; C₂-C₂₀ alkoxyalkyl; C₂-C₂₀ alkylthioalkyl; C₂-C₂₀ alkylsulfinylalkyl; C₂-C₂₀ alkylsulfonylalkyl; C₅-C₂₀ cycloalkylalkyl; C₄-C₂₀ alkenyloxyalkyl; C₄-C₂₀ alkynyloxyalkyl; C₄-C₂₀ (cycloalkyl) oxyalkyl; C₄-C₂₀ alkenylthioalkyl; C₄-C₂₀ alkynylthioalkyl; C₆-C₂₀ (cycloalkyl) thioalkyl; C₂-C₂₀ haloalkoxyalkyl; C₄-C₂₀ haloalkenyloxyalkyl; C₄-C₂₀ haloalkynyloxyalkyl; C₄-C₂₀ alkoxyalkenyl; C₄-C₂₀ alkoxyalkynyl; C₄-C₂₀ alkylthioalkenyl; C₄-C₂₀ alkylthioalkynyl; C₄-C₂₀ trialkylsilylalkyl; C₁-C₂₀ alkyl substituted with NR₃R₄, nitro, cyano, or phenyl optionally substituted with R₅, R₆, and R₇; C₁-C₂₀ alkoxy; C₁-C₂₀ haloalkoxy; C₁-C₂₀ alkylthio; C₁-C₂₀ haloalkylthio; NR₃R₄; or phenyl, benzyl, pyridyl, furanyl, thienyl, naphthyl, pyrimidinyl, benzofuranyl, benzothienyl, or quinolinyl each optionally substituted with R₅, R₆ or R₇;

R₂ is C₁-C₂₀ alkyl; C₃-C₆ cycloalkyl; C₄-C₂₀ alkenyl; C₄-C₂₀ alkynyl; C₁-C₂₀ haloalkyl; C₃-C₂₀ haloalkenyl; C₃-C₂₀ haloalkynyl; C₂-C₂₀ alkoxyalkyl; C₂-C₂₀ alkylthioalkyl; C₂-C₂₀ alkylsulfinylalkyl; C₂-C₂₀ alkylsulfonylalkyl; C₅-C₂₀ cycloalkylalkyl; C₄-C₂₀ alkenyloxyalkyl; C₄-C₂₀ alkynyloxyalkyl; C₄-C₂₀ (cycloalkyl) oxyalkyl; C₄-C₂₀ alkenylthioalkyl; C₄-C₂₀ alkynylthioalkyl; C₆-C₂₀ (cycloalkyl) thioalkyl; C₂-C₂₀ haloalkoxyalkyl; C₄-C₂₀ haloalkenyloxyalkyl; C₄-C₂₀ haloalkynyloxyalkyl; C₄-C₂₀ alkoxyalkenyl; C₄-C₂₀ alkoxyalkynyl; C₄-C₂₀ alkylthioalkenyl; C₄-C₂₀ alkylthioalkynyl; C₄-C₂₀ trialkylsilylalkyl; C₁-C₂₀ alkyl substituted with NR₃R₄, nitro, cyano, or phenyl optionally substituted with R₅, R₆, and R₇; C₁-C₂₀ alkoxy; C₁-C₂₀ haloalkoxy; C₁-C₂₀ alkylthio; C₁-C₂₀ haloalkylthio; NR₃R₄; or phenyl, benzyl, pyridyl, furanyl, thienyl, naphthyl, pyrimidinyl, benzofuranyl, benzothienyl, or

quinolinyll each optionally substituted with R₅, R₆ or R₇;

R₃ is independently hydrogen; C₁-C₄ alkyl; or phenyl optionally substituted with at least one R₈;

R₄ is independently hydrogen; C₁-C₈ alkyl; or phenyl optionally substituted with at least one R₈;

R₅ is independently C₁-C₆ alkyl; C₁-C₆ alkoxy; C₁-C₆ haloalkyl; halogen; C₂-C₈ alkynyl; C₁-C₆ thioalkyl; phenyl or phenoxy each optionally substituted with at least one R₈; cyano; nitro; C₁-C₆ haloalkoxy; C₁-C₆ haloalkythio; C₂-C₆ alkenyl; C₂-C₆ haloalkenyl; acetyl; CO₂CH₃; or N(C₁-C₂ alkyl)₂;

R₆ is independently methyl; ethyl; methoxy; methylthio; halogen; or trifluoromethyl;

R₇ is independently halogen; and

R₈ is independently halogen; C₁-C₄ alkyl; C₁-C₄ alkoxy; C₁-C₄ haloalkyl; nitro; or cyano;

wherein:

the amino acid in the first position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, lysine, methionine, serine, threonine and tryptophan;

the amino acid in the second position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, histidine, cysteine, threonine, tyrosine, and tryptophan; and

the amino acids in positions three through six, based on numbered amino acids from N-terminus to C-terminus, are any amino acid.

10. The antimicrobial composition of claim 9 wherein the amino acids in the first and second positions of said peptide, based on numbered amino acids from N-terminus to C-terminus, are selected from the group consisting of Arg-Tyr, Arg-Cys, Ser-Thr, Met-Trp, Lys-Trp, Thr-Trp, Trp-Arg, Trp-His, and Trp-Tyr.

11. The antimicrobial composition of claim 9 wherein said peptides are incorporated into a

polymer.

12. The antimicrobial composition of claim 11 wherein said polymer is selected from the group consisting of a polysaccharide, a glycol polymer, a polyester, a polyurethane, a polyacrylate, a polyacrylonitrile, a polyamide, a polyolefin, a polystyrene, a vinyl polymer, a polypropylene, silk, a biopolymer, and mixtures thereof.

13. An antimicrobial composition comprising a plurality of hexapeptides and at least one carrier, wherein for each hexapeptide:

the amino acid in the first position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, lysine, methionine, serine, threonine and tryptophan;

the amino acid in the second position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, histidine, cysteine, threonine, tyrosine, and tryptophan;

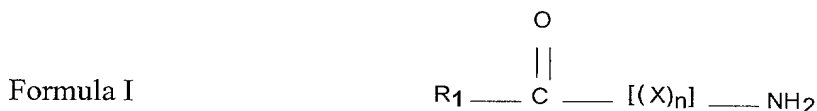
the amino acids in positions three through six, based on numbered amino acids from N-terminus to C-terminus, are any amino acid; and

wherein the first two amino acids of said hexapeptides are other than arginine-arginine, tryptophan-tryptophan, tryptophan-cysteine, tryptophan-lysine, arginine-tryptophan, or threonine-arginine.

14. The antimicrobial composition of claim 13 wherein the amino acids in the first and second positions of said peptides, based on numbered amino acids from N-terminus to C-terminus, are selected from the group consisting of Arg-Tyr, Arg-Cys, Ser-Thr, Met-Trp, Lys-Trp, Thr-Trp, Trp-Arg, Trp-His, and Trp-Tyr.

15. The antimicrobial composition of claim 13 wherein said carrier is selected from the group consisting of a pharmaceutically acceptable carrier, an industrially acceptable carrier, a household product, and a personal care composition.

16. An antimicrobial composition comprising a plurality of hexapeptides and at least one carrier, wherein said each hexapeptide is represented by Formula I:



wherein:

X represents any amino acid except glutamate or aspartate;

n = 6;

R₁ is C₁-C₂₀ alkyl; C₃-C₆ cycloalkyl; C₄-C₂₀ alkenyl; C₄-C₂₀ alkynyl; C₁-C₂₀ haloalkyl; C₃-C₂₀ haloalkenyl; C₃-C₂₀ haloalkynyl; C₂-C₂₀ alkoxyalkyl; C₂-C₂₀ alkylthioalkyl; C₂-C₂₀ alkylsulfinylalkyl; C₂-C₂₀ alkylsulfonylalkyl; C₅-C₂₀ cycloalkylalkyl; C₄-C₂₀ alkenyloxyalkyl; C₄-C₂₀ alkynyloxyalkyl; C₄-C₂₀ (cycloalkyl) oxyalkyl; C₄-C₂₀ alkenylthioalkyl; C₄-C₂₀ alkynylthioalkyl; C₆-C₂₀ (cycloalkyl) thioalkyl; C₂-C₂₀ haloalkoxyalkyl; C₄-C₂₀ haloalkenyloxyalkyl; C₄-C₂₀ haloalkynyloxyalkyl; C₄-C₂₀ alkoxyalkenyl; C₄-C₂₀ alkoxyalkynyl; C₄-C₂₀ alkylthioalkenyl; C₄-C₂₀ alkylthioalkynyl; C₄-C₂₀ trialkylsilylalkyl; C₁-C₂₀ alkyl substituted with NR₃R₄, nitro, cyano, or phenyl optionally substituted with R₅, R₆, and R₇; C₁-C₂₀ alkoxy; C₁-C₂₀ haloalkoxy; C₁-C₂₀ alkylthio; C₁-C₂₀ haloalkylthio; NR₃R₄; or phenyl, benzyl, pyridyl, furanyl, thienyl, naphthyl, pyrimidinyl, benzofuranyl, benzothienyl, or quinolinyl each optionally substituted with R₅, R₆ or R₇;

R₃ is independently hydrogen; C₁-C₄ alkyl; or phenyl optionally substituted with at least one R₈;

R₄ is independently hydrogen; C₁-C₈ alkyl; or phenyl optionally substituted with at least one R₈;

R₅ is independently C₁-C₆ alkyl; C₁-C₆ alkoxy; C₁-C₆ haloalkyl; halogen; C₂-C₈ alkynyl; C₁-C₆ thioalkyl; phenyl or phenoxy each optionally substituted with at least one R₈; cyano; nitro; C₁-C₆ haloalkoxy; C₁-C₆ haloalkylthio; C₂-C₆ alkenyl; C₂-C₆ haloalkenyl; acetyl; CO₂CH₃; or N(C₁-C₂ alkyl)₂;

R₆ is independently methyl; ethyl; methoxy; methylthio; halogen; or trifluoromethyl;
and

R₇ is independently halogen;

R₈ is independently halogen; C₁-C₄ alkyl; C₁-C₄ alkoxy; C₁-C₄ haloalkyl; nitro; or
cyano;
wherein:

the amino acid in the first position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, lysine, methionine, serine, threonine and tryptophan;

the amino acid in the second position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, histidine, cysteine, threonine, tyrosine, and tryptophan; and

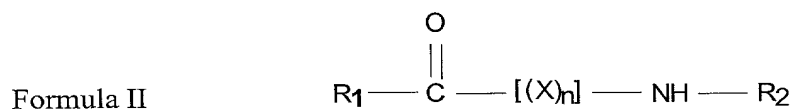
the amino acids in positions three through six, based on numbered amino acids from N-terminus to C-terminus, are any amino acid;

wherein the first two amino acids of said hexapeptides are other than arginine-arginine, tryptophan-tryptophan, tryptophan-cysteine, tryptophan-lysine, arginine-tryptophan, or threonine-arginine.

17. The antimicrobial composition of claim 16 wherein the amino acids in the first and second positions of said peptides, based on numbered amino acids from N-terminus to C-terminus, are selected from the group consisting of Arg-Tyr, Arg-Cys, Ser-Thr, Met-Trp, Lys-Trp, Thr-Trp, Trp-Arg, Trp-His, and Trp-Tyr.

18. The antimicrobial composition of claim 16 wherein said carrier is selected from the group consisting of a pharmaceutically acceptable carrier, an industrially acceptable carrier, a household product, and a personal care composition.

19. An antimicrobial composition comprising a plurality of hexapeptides and at least one carrier, wherein said each hexapeptide is represented by Formula II:



wherein:

X represents any amino acid except glutamate or aspartate;

n = 6;

R₁ is C₁-C₂₀ alkyl; C₃-C₆ cycloalkyl; C₄-C₂₀ alkenyl; C₄-C₂₀ alkynyl; C₁-C₂₀ haloalkyl; C₃-C₂₀ haloalkenyl; C₃-C₂₀ haloalkynyl; C₂-C₂₀ alkoxyalkyl; C₂-C₂₀ alkylthioalkyl; C₂-C₂₀ alkylsulfanylalkyl; C₂-C₂₀ alkylsulfonylalkyl; C₅-C₂₀ cycloalkylalkyl; C₄-C₂₀ alkenyloxyalkyl; C₄-C₂₀ alkynyloxyalkyl; C₄-C₂₀ (cycloalkyl) oxyalkyl; C₄-C₂₀ alkenylthioalkyl; C₄-C₂₀ alkynylthioalkyl; C₆-C₂₀ (cycloalkyl) thioalkyl; C₂-C₂₀ haloalkoxyalkyl; C₄-C₂₀ haloalkenyloxyalkyl; C₄-C₂₀ haloalkynyloxyalkyl; C₄-C₂₀ alkoxyalkenyl; C₄-C₂₀ alkoxyalkynyl; C₄-C₂₀ alkylthioalkenyl; C₄-C₂₀ alkylthioalkynyl; C₄-C₂₀ trialkylsilylalkyl; C₁-C₂₀ alkyl substituted with NR₃R₄, nitro, cyano, or phenyl optionally substituted with R₅, R₆, and R₇; C₁-C₂₀ alkoxy; C₁-C₂₀ haloalkoxy; C₁-C₂₀ alkylthio; C₁-C₂₀ haloalkylthio; NR₃R₄; or phenyl, benzyl, pyridyl, furanyl, thienyl, naphthyl, pyrimidinyl, benzofuranyl, benzothienyl, or quinolinyl each optionally substituted with R₅, R₆ or R₇;

R₂ is C₁-C₂₀ alkyl; C₃-C₆ cycloalkyl; C₄-C₂₀ alkenyl; C₄-C₂₀ alkynyl; C₁-C₂₀ haloalkyl; C₃-C₂₀ haloalkenyl; C₃-C₂₀ haloalkynyl; C₂-C₂₀ alkoxyalkyl; C₂-C₂₀ alkylthioalkyl; C₂-C₂₀ alkylsulfanylalkyl; C₂-C₂₀ alkylsulfonylalkyl; C₅-C₂₀ cycloalkylalkyl; C₄-C₂₀ alkenyloxyalkyl; C₄-C₂₀ alkynyloxyalkyl; C₄-C₂₀ (cycloalkyl) oxyalkyl; C₄-C₂₀ alkenylthioalkyl; C₄-C₂₀ alkynylthioalkyl; C₆-C₂₀ (cycloalkyl) thioalkyl; C₂-C₂₀ haloalkoxyalkyl; C₄-C₂₀ haloalkenyloxyalkyl; C₄-C₂₀ haloalkynyloxyalkyl; C₄-C₂₀ alkoxyalkenyl; C₄-C₂₀ alkoxyalkynyl; C₄-C₂₀ alkylthioalkenyl; C₄-C₂₀ alkylthioalkynyl; C₄-C₂₀ trialkylsilylalkyl; C₁-C₂₀ alkyl substituted with NR₃R₄, nitro, cyano, or phenyl optionally substituted with R₅, R₆, and R₇; C₁-C₂₀ alkoxy; C₁-C₂₀ haloalkoxy; C₁-C₂₀ alkylthio; C₁-C₂₀ haloalkylthio; NR₃R₄; or phenyl, benzyl, pyridyl, furanyl, thienyl, naphthyl, pyrimidinyl, benzofuranyl, benzothienyl, or quinolinyl each optionally substituted with R₅, R₆ or R₇;

R₃ is independently hydrogen; C₁-C₄ alkyl; or phenyl optionally substituted with at least one R₈;

R₄ is independently hydrogen; C₁-C₈ alkyl; or phenyl optionally substituted with at least one R₈;

R₅ is independently C₁-C₆ alkyl; C₁-C₆ alkoxy; C₁-C₆ haloalkyl; halogen; C₂-C₈ alkynyl; C₁-C₆ thioalkyl; phenyl or phenoxy each optionally substituted with at least one R₈; cyano; nitro; C₁-C₆ haloalkoxy; C₁-C₆ haloalkythio; C₂-C₆ alkenyl; C₂-C₆ haloalkenyl; acetyl; CO₂CH₃; or N(C₁-C₂ alkyl)₂;

R₆ is independently methyl; ethyl; methoxy; methylthio; halogen; or trifluoromethyl;

R₇ is independently halogen; and

R₈ is independently halogen; C₁-C₄ alkyl; C₁-C₄ alkoxy; C₁-C₄ haloalkyl; nitro; or cyano.

20. The antimicrobial composition of claim 19 wherein the amino acids in the first and second positions, based on numbered amino acids from N-terminus to C-terminus, are selected from the group consisting of Arg-Tyr, Arg-Cys, Ser-Thr, Met-Trp, Lys-Trp, Thr-Trp, Trp-Arg, Trp-His, and Trp-Tyr.

21. The antimicrobial composition of claim 19 wherein said carrier is selected from the group consisting of a pharmaceutically acceptable carrier, an industrially acceptable carrier, a household product, and a personal care composition.

22. A method for preventing, inhibiting, or terminating the growth of at least one microbe comprising administering an antimicrobial amount of a plurality of hexapeptides and at least one carrier, wherein for each hexapeptide:

the amino acid in the first position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, lysine, methionine, serine, threonine and tryptophan;

the amino acid in the second position, based on numbered amino acids from N-

terminus to C-terminus, is selected from the group consisting of arginine, histidine, cysteine, threonine, tyrosine, and tryptophan;

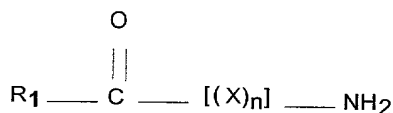
the amino acids in positions three through six, based on numbered amino acids from N-terminus to C-terminus, are any amino acid; and

wherein the first two amino acids of said hexapeptides are other than arginine-arginine, tryptophan-tryptophan, tryptophan-cysteine, tryptophan-lysine, arginine-tryptophan, or threonine-arginine.

23. The method of claim 22 wherein said microbe comprises *Burkholderia cepacia*.

24. A method for preventing, inhibiting, or terminating the growth of at least one microbe comprising administering an antimicrobial amount of a plurality of peptides and at least one carrier, wherein said peptides are each represented by Formula I:

Formula I



wherein:

X represents any amino acid except glutamate or aspartate;

n = 6;

R₁ is C₁-C₂₀ alkyl; C₃-C₆ cycloalkyl; C₄-C₂₀ alkenyl; C₄-C₂₀ alkynyl; C₁-C₂₀ haloalkyl; C₃-C₂₀ haloalkenyl; C₃-C₂₀ haloalkynyl; C₂-C₂₀ alkoxyalkyl; C₂-C₂₀ alkylthioalkyl; C₂-C₂₀ alkylsulfinylalkyl; C₂-C₂₀ alkylsulfonylalkyl; C₅-C₂₀ cycloalkylalkyl; C₄-C₂₀ alkenyloxyalkyl; C₄-C₂₀ alkynyloxyalkyl; C₄-C₂₀ (cycloalkyl) oxyalkyl; C₄-C₂₀ alkenylthioalkyl; C₄-C₂₀ alkynylthioalkyl; C₆-C₂₀ (cycloalkyl) thioalkyl; C₂-C₂₀ haloalkoxyalkyl; C₄-C₂₀ haloalkenyloxyalkyl; C₄-C₂₀ haloalkynyloxyalkyl; C₄-C₂₀ alkoxyalkenyl; C₄-C₂₀ alkoxyalkynyl; C₄-C₂₀ alkylthioalkenyl; C₄-C₂₀ alkylthioalkynyl; C₄-C₂₀ trialkylsilylalkyl; C₁-C₂₀ alkyl substituted with NR₃R₄, nitro, cyano, or phenyl optionally substituted with R₅, R₆, and R₇; C₁-C₂₀ alkoxy; C₁-C₂₀ haloalkoxy; C₁-C₂₀ alkylthio; C₁-C₂₀ haloalkylthio; NR₃R₄; or phenyl, benzyl, pyridyl, furanyl, thienyl, naphthyl, pyrimidinyl, benzofuranyl, benzothienyl, or

quinolinyl each optionally substituted with R₅, R₆ or R₇;

R₂ is C₁-C₂₀ alkyl; C₃-C₆ cycloalkyl; C₄-C₂₀ alkenyl; C₄-C₂₀ alkynyl; C₁-C₂₀ haloalkyl; C₃-C₂₀ haloalkenyl; C₃-C₂₀ haloalkynyl; C₂-C₂₀ alkoxyalkyl; C₂-C₂₀ alkylthioalkyl; C₂-C₂₀ alkylsulfanylalkyl; C₂-C₂₀ alkylsulfonylalkyl; C₅-C₂₀ cycloalkylalkyl; C₄-C₂₀ alkenyloxyalkyl; C₄-C₂₀ alkynyloxyalkyl; C₄-C₂₀ (cycloalkyl) oxyalkyl; C₄-C₂₀ alkenylthioalkyl; C₄-C₂₀ alkynylthioalkyl; C₆-C₂₀ (cycloalkyl) thioalkyl; C₂-C₂₀ haloalkoxyalkyl; C₄-C₂₀ haloalkenyloxyalkyl; C₄-C₂₀ haloalkynyloxyalkyl; C₄-C₂₀ alkoxyalkenyl; C₄-C₂₀ alkoxyalkynyl; C₄-C₂₀ alkylthioalkenyl; C₄-C₂₀ alkylthioalkynyl; C₄-C₂₀ trialkylsilylalkyl; C₁-C₂₀ alkyl substituted with NR₃R₄, nitro, cyano, or phenyl optionally substituted with R₅, R₆, and R₇; C₁-C₂₀ alkoxy; C₁-C₂₀ haloalkoxy; C₁-C₂₀ alkylthio; C₁-C₂₀ haloalkylthio; NR₃R₄; or phenyl, benzyl, pyridyl, furanyl, thienyl, naphthyl, pyrimidinyl, benzofuranyl, benzothienyl, or quinolinyl each optionally substituted with R₅, R₆ or R₇;

R₃ is independently hydrogen; C₁-C₄ alkyl; or phenyl optionally substituted with at least one R₈;

R₄ is independently hydrogen; C₁-C₈ alkyl; or phenyl optionally substituted with at least one R₈;

R₅ is independently C₁-C₆ alkyl; C₁-C₆ alkoxy; C₁-C₆ haloalkyl; halogen; C₂-C₈ alkynyl; C₁-C₆ thioalkyl; phenyl or phenoxy each optionally substituted with at least one R₈; cyano; nitro; C₁-C₆ haloalkoxy; C₁-C₆ haloalkylthio; C₂-C₆ alkenyl; C₂-C₆ haloalkenyl; acetyl; CO₂CH₃; or N(C₁-C₂ alkyl)₂;

R₆ is independently methyl; ethyl; methoxy; methylthio; halogen; or trifluoromethyl;

R₇ is independently halogen; and

R₈ is independently halogen; C₁-C₄ alkyl; C₁-C₄ alkoxy; C₁-C₄ haloalkyl; nitro; or cyano;

wherein:

the amino acid in the first position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, lysine, methionine, serine, threonine and tryptophan;

the amino acid in the second position, based on numbered amino acids from N-

terminus to C-terminus, is selected from the group consisting of arginine, histidine, cysteine, threonine, tyrosine, and tryptophan; and

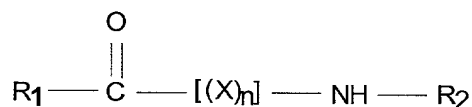
the amino acids in positions three through six, based on numbered amino acids from N-terminus to C-terminus, are any amino acid;

wherein the first two amino acids of said hexapeptides are other than arginine-arginine, tryptophan-tryptophan, tryptophan-cysteine, tryptophan-lysine, arginine-tryptophan, or threonine-arginine.

25. The method of claim 24 wherein said microbe comprises *Burkholderia cepacia*.

26. A method for preventing, inhibiting, or terminating the growth of at least one microbe comprising administering an antimicrobial amount of a plurality of peptides and at least one carrier, wherein said peptides are each represented by Formula II:

Formula II



wherein:

X represents any amino acid except glutamate or aspartate;

n = 6;

R₁ is C₁-C₂₀ alkyl; C₃-C₆ cycloalkyl; C₄-C₂₀ alkenyl; C₄-C₂₀ alkynyl; C₁-C₂₀ haloalkyl; C₃-C₂₀ haloalkenyl; C₃-C₂₀ haloalkynyl; C₂-C₂₀ alkoxyalkyl; C₂-C₂₀ alkylthioalkyl; C₂-C₂₀ alkylsulfinylalkyl; C₂-C₂₀ alkylsulfonylalkyl; C₅-C₂₀ cycloalkylalkyl; C₄-C₂₀ alkenyloxyalkyl; C₄-C₂₀ alkynyloxyalkyl; C₄-C₂₀ (cycloalkyl) oxyalkyl; C₄-C₂₀ alkenylthioalkyl; C₄-C₂₀ alkynylthioalkyl; C₆-C₂₀ (cycloalkyl) thioalkyl; C₂-C₂₀ haloalkoxyalkyl; C₄-C₂₀ haloalkenyloxyalkyl; C₄-C₂₀ haloalkynyloxyalkyl; C₄-C₂₀ alkoxyalkenyl; C₄-C₂₀ alkoxyalkynyl; C₄-C₂₀ alkylthioalkenyl; C₄-C₂₀ alkylthioalkynyl; C₄-C₂₀ trialkylsilylalkyl; C₁-C₂₀ alkyl substituted with NR₃R₄, nitro, cyano, or phenyl optionally substituted with R₅, R₆, and R₇; C₁-C₂₀ alkoxy; C₁-C₂₀ haloalkoxy; C₁-C₂₀ alkylthio; C₁-C₂₀ haloalkylthio; NR₃R₄; or phenyl, benzyl, pyridyl, furanyl, thienyl, naphthyl, pyrimidinyl, benzofuranyl, benzothienyl, or

quinoliny each optionally substituted with R₅, R₆ or R₇;

R₂ is C₁-C₂₀ alkyl; C₃-C₆ cycloalkyl; C₄-C₂₀ alkenyl; C₄-C₂₀ alkynyl; C₁-C₂₀ haloalkyl; C₃-C₂₀ haloalkenyl; C₃-C₂₀ haloalkynyl; C₂-C₂₀ alkoxyalkyl; C₂-C₂₀ alkylthioalkyl; C₂-C₂₀ alkylsulfinylalkyl; C₂-C₂₀ alkylsulfonylalkyl; C₅-C₂₀ cycloalkylalkyl; C₄-C₂₀ alkenyloxyalkyl; C₄-C₂₀ alkynyloxyalkyl; C₄-C₂₀ (cycloalkyl) oxyalkyl; C₄-C₂₀ alkenylthioalkyl; C₄-C₂₀ alkynylthioalkyl; C₆-C₂₀ (cycloalkyl) thioalkyl; C₂-C₂₀ haloalkoxyalkyl; C₄-C₂₀ haloalkenyloxyalkyl; C₄-C₂₀ haloalkynyloxyalkyl; C₄-C₂₀ alkoxyalkenyl; C₄-C₂₀ alkoxyalkynyl; C₄-C₂₀ alkylthioalkenyl; C₄-C₂₀ alkylthioalkynyl; C₄-C₂₀ trialkylsilylalkyl; C₁-C₂₀ alkyl substituted with NR₃R₄, nitro, cyano, or phenyl optionally substituted with R₅, R₆, and R₇; C₁-C₂₀ alkoxy; C₁-C₂₀ haloalkoxy; C₁-C₂₀ alkylthio; C₁-C₂₀ haloalkylthio; NR₃R₄; or phenyl, benzyl, pyridyl, furanyl, thienyl, naphthyl, pyrimidinyl, benzofuranyl, benzothienyl, or quinoliny each optionally substituted with R₅, R₆ or R₇;

R₃ is independently hydrogen; C₁-C₄ alkyl; or phenyl optionally substituted with at least one R₈;

R₄ is independently hydrogen; C₁-C₈ alkyl; or phenyl optionally substituted with at least one R₈;

R₅ is independently C₁-C₆ alkyl; C₁-C₆ alkoxy; C₁-C₆ haloalkyl; halogen; C₂-C₈ alkynyl; C₁-C₆ thioalkyl; phenyl or phenoxy each optionally substituted with at least one R₈; cyano; nitro; C₁-C₆ haloalkoxy; C₁-C₆ haloalkylthio; C₂-C₆ alkenyl; C₂-C₆ haloalkenyl; acetyl; CO₂CH₃; or N(C₁-C₂ alkyl)₂;

R₆ is independently methyl; ethyl; methoxy; methylthio; halogen; or trifluoromethyl;

R₇ is independently halogen; and

R₈ is independently halogen; C₁-C₄ alkyl; C₁-C₄ alkoxy; C₁-C₄ haloalkyl; nitro; or cyano.

27. The method of claim 26 wherein said microbe comprises *Burkholderia cepacia*.

28. A composition for coating a substrate comprising an antimicrobial amount of a plurality of hexapeptides and at least one carrier, wherein for each hexapeptide:

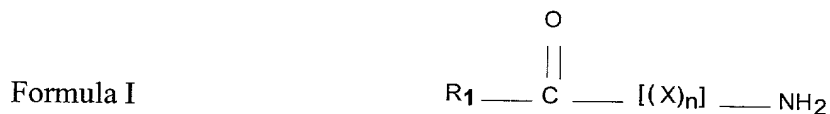
the amino acid in the first position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, lysine, methionine, serine, threonine and tryptophan;

the amino acid in the second position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, histidine, cysteine, threonine, tyrosine, and tryptophan;

the amino acids in positions three through six, based on numbered amino acids from N-terminus to C-terminus, are any amino acid; and

wherein the first two amino acids of said hexapeptides are other than arginine-arginine, tryptophan-tryptophan, tryptophan-cysteine, tryptophan-lysine, arginine-tryptophan, or threonine-arginine.

29. A composition for coating a substrate comprising an antimicrobial amount of a plurality of peptides and at least one carrier, wherein each of said peptides are represented by Formula I:



wherein:

X represents any amino acid except glutamate or aspartate;

n = 6;

R₁ is C₁-C₂₀ alkyl; C₃-C₆ cycloalkyl; C₄-C₂₀ alkenyl; C₄-C₂₀ alkynyl; C₁-C₂₀ haloalkyl; C₃-C₂₀ haloalkenyl; C₃-C₂₀ haloalkynyl; C₂-C₂₀ alkoxyalkyl; C₂-C₂₀ alkylthioalkyl; C₂-C₂₀ alkylsulfinylalkyl; C₂-C₂₀ alkylsulfonylalkyl; C₅-C₂₀ cycloalkylalkyl; C₄-C₂₀ alkenyloxyalkyl; C₄-C₂₀ alkynyloxyalkyl; C₄-C₂₀ (cycloalkyl) oxyalkyl; C₄-C₂₀ alkenylthioalkyl; C₄-C₂₀ alkynylthioalkyl; C₆-C₂₀ (cycloalkyl) thioalkyl; C₂-C₂₀ haloalkoxyalkyl; C₄-C₂₀ haloalkenyloxyalkyl; C₄-C₂₀ haloalkynyloxyalkyl; C₄-C₂₀ alkoxyalkenyl; C₄-C₂₀ alkoxyalkynyl; C₄-C₂₀ alkylthioalkenyl; C₄-C₂₀ alkylthioalkynyl; C₄-C₂₀ trialkylsilylalkyl; C₁-

C₂₀ alkyl substituted with NR₃R₄, nitro, cyano, or phenyl optionally substituted with R₅, R₆, and R₇; C₁-C₂₀ alkoxy; C₁-C₂₀ haloalkoxy; C₁-C₂₀ alkylthio; C₁-C₂₀ haloalkylthio; NR₃R₄; or phenyl, benzyl, pyridyl, furanyl, thienyl, naphthyl, pyrimidinyl, benzofuranyl, benzothienyl, or quinolinyl each optionally substituted with R₅, R₆ or R₇;

R₂ is C₁-C₂₀ alkyl; C₃-C₆ cycloalkyl; C₄-C₂₀ alkenyl; C₄-C₂₀ alkynyl; C₁-C₂₀ haloalkyl; C₃-C₂₀ haloalkenyl; C₃-C₂₀ haloalkynyl; C₂-C₂₀ alkoxyalkyl; C₂-C₂₀ alkylthioalkyl; C₂-C₂₀ alkylsulfanylalkyl; C₂-C₂₀ alkylsulfonylalkyl; C₅-C₂₀ cycloalkylalkyl; C₄-C₂₀ alkenyloxyalkyl; C₄-C₂₀ alkynyloxyalkyl; C₄-C₂₀ (cycloalkyl) oxyalkyl; C₄-C₂₀ alkenylthioalkyl; C₄-C₂₀ alkynylthioalkyl; C₆-C₂₀ (cycloalkyl) thioalkyl; C₂-C₂₀ haloalkoxyalkyl; C₄-C₂₀ haloalkenyloxyalkyl; C₄-C₂₀ haloalkynyloxyalkyl; C₄-C₂₀ alkoxyalkenyl; C₄-C₂₀ alkoxyalkynyl; C₄-C₂₀ alkylthioalkenyl; C₄-C₂₀ alkylthioalkynyl; C₄-C₂₀ trialkylsilylalkyl; C₁-C₂₀ alkyl substituted with NR₃R₄, nitro, cyano, or phenyl optionally substituted with R₅, R₆, and R₇; C₁-C₂₀ alkoxy; C₁-C₂₀ haloalkoxy; C₁-C₂₀ alkylthio; C₁-C₂₀ haloalkylthio; NR₃R₄; or phenyl, benzyl, pyridyl, furanyl, thienyl, naphthyl, pyrimidinyl, benzofuranyl, benzothienyl, or quinolinyl each optionally substituted with R₅, R₆ or R₇;

R₃ is independently hydrogen; C₁-C₄ alkyl; or phenyl optionally substituted with at least one R₈;

R₄ is independently hydrogen; C₁-C₈ alkyl; or phenyl optionally substituted with at least one R₈;

R₅ is independently C₁-C₆ alkyl; C₁-C₆ alkoxy; C₁-C₆ haloalkyl; halogen; C₂-C₈ alkynyl; C₁-C₆ thioalkyl; phenyl or phenoxy each optionally substituted with at least one R₈; cyano; nitro; C₁-C₆ haloalkoxy; C₁-C₆ haloalkylthio; C₂-C₆ alkenyl; C₂-C₆ haloalkenyl; acetyl; CO₂CH₃; or N(C₁-C₂ alkyl)₂;

R₆ is independently methyl; ethyl; methoxy; methylthio; halogen; or trifluoromethyl;

R₇ is independently halogen; and

R₈ is independently halogen; C₁-C₄ alkyl; C₁-C₄ alkoxy; C₁-C₄ haloalkyl; nitro; or cyano;

wherein:

the amino acid in the first position, based on numbered amino acids from N-terminus to

C-terminus, is selected from the group consisting of arginine, lysine, methionine, serine, threonine and tryptophan;

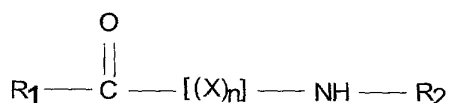
the amino acid in the second position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, histidine, cysteine, threonine, tyrosine, and tryptophan; and

the amino acids in positions three through six, based on numbered amino acids from N-terminus to C-terminus, are any amino acid;

wherein the first two amino acids of said hexapeptides are other than arginine-arginine, tryptophan-tryptophan, tryptophan-cysteine, tryptophan-lysine, arginine-tryptophan, or threonine-arginine.

30. A composition for coating a substrate comprising an antimicrobial amount of a plurality of peptides and at least one carrier, wherein each of said peptides are represented by Formula II:

Formula II



wherein:

X represents any amino acid except glutamate or aspartate;

n = 6;

R₁ is C₁-C₂₀ alkyl; C₃-C₆ cycloalkyl; C₄-C₂₀ alkenyl; C₄-C₂₀ alkynyl; C₁-C₂₀ haloalkyl; C₃-C₂₀ haloalkenyl; C₃-C₂₀ haloalkynyl; C₂-C₂₀ alkoxyalkyl; C₂-C₂₀ alkylthioalkyl; C₂-C₂₀ alkylsulfinylalkyl; C₂-C₂₀ alkylsulfonylalkyl; C₅-C₂₀ cycloalkylalkyl; C₄-C₂₀ alkenyloxyalkyl; C₄-C₂₀ alkynyloxyalkyl; C₄-C₂₀ (cycloalkyl) oxyalkyl; C₄-C₂₀ alkenylthioalkyl; C₄-C₂₀ alkynylthioalkyl; C₆-C₂₀ (cycloalkyl) thioalkyl; C₂-C₂₀ haloalkoxyalkyl; C₄-C₂₀ haloalkenyloxyalkyl; C₄-C₂₀ haloalkynyloxyalkyl; C₄-C₂₀ alkoxyalkenyl; C₄-C₂₀ alkoxyalkynyl; C₄-C₂₀ alkylthioalkenyl; C₄-C₂₀ alkylthioalkynyl; C₄-C₂₀ trialkylsilylalkyl; C₁-C₂₀ alkyl substituted with NR₃R₄, nitro, cyano, or phenyl optionally substituted with R₅, R₆, and R₇; C₁-C₂₀ alkoxy; C₁-C₂₀ haloalkoxy; C₁-C₂₀ alkylthio; C₁-C₂₀ haloalkylthio; NR₃R₄; or

phenyl, benzyl, pyridyl, furanyl, thienyl, naphthyl, pyrimidinyl, benzofuranyl, benzothienyl, or quinolinyl each optionally substituted with R₅, R₆ or R₇;

R₂ is C₁-C₂₀ alkyl; C₃-C₆ cycloalkyl; C₄-C₂₀ alkenyl; C₄-C₂₀ alkynyl; C₁-C₂₀ haloalkyl; C₃-C₂₀ haloalkenyl; C₃-C₂₀ haloalkynyl; C₂-C₂₀ alkoxyalkyl; C₂-C₂₀ alkylthioalkyl; C₂-C₂₀ alkylsulfinylalkyl; C₂-C₂₀ alkylsulfonylalkyl; C₅-C₂₀ cycloalkylalkyl; C₄-C₂₀ alkenyloxyalkyl; C₄-C₂₀ alkynyloxyalkyl; C₄-C₂₀ (cycloalkyl) oxyalkyl; C₄-C₂₀ alkenylthioalkyl; C₄-C₂₀ alkynylthioalkyl; C₆-C₂₀ (cycloalkyl) thioalkyl; C₂-C₂₀ haloalkoxyalkyl; C₄-C₂₀ haloalkenyloxyalkyl; C₄-C₂₀ haloalkynyloxyalkyl; C₄-C₂₀ alkoxyalkenyl; C₄-C₂₀ alkoxyalkynyl; C₄-C₂₀ alkylthioalkenyl; C₄-C₂₀ alkylthioalkynyl; C₄-C₂₀ trialkylsilylalkyl; C₁-C₂₀ alkyl substituted with NR₃R₄, nitro, cyano, or phenyl optionally substituted with R₅, R₆, and R₇; C₁-C₂₀ alkoxy; C₁-C₂₀ haloalkoxy; C₁-C₂₀ alkylthio; C₁-C₂₀ haloalkylthio; NR₃R₄; or phenyl, benzyl, pyridyl, furanyl, thienyl, naphthyl, pyrimidinyl, benzofuranyl, benzothienyl, or quinolinyl each optionally substituted with R₅, R₆ or R₇;

R₃ is independently hydrogen; C₁-C₄ alkyl; or phenyl optionally substituted with at least one R₈;

R₄ is independently hydrogen; C₁-C₈ alkyl; or phenyl optionally substituted with at least one R₈;

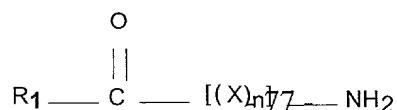
R₅ is independently C₁-C₆ alkyl; C₁-C₆ alkoxy; C₁-C₆ haloalkyl; halogen; C₂-C₈ alkynyl; C₁-C₆ thioalkyl; phenyl or phenoxy each optionally substituted with at least one R₈; cyano; nitro; C₁-C₆ haloalkoxy; C₁-C₆ haloalkylthio; C₂-C₆ alkenyl; C₂-C₆ haloalkenyl; acetyl; CO₂CH₃; or N(C₁-C₂ alkyl)₂;

R₆ is independently methyl; ethyl; methoxy; methylthio; halogen; or trifluoromethyl;

R₇ is independently halogen; and

R₈ is independently halogen; C₁-C₄ alkyl; C₁-C₄ alkoxy; C₁-C₄ haloalkyl; nitro; or cyano.

31. An antimicrobial composition comprising a plurality of peptides, wherein said peptides each are represented by Formula I:



Formula I

wherein:

X represents any amino acid except glutamate or aspartate;

n = 1-10;

R₁ is C₁-C₂₀ alkyl; C₃-C₆ cycloalkyl; C₄-C₂₀ alkenyl; C₄-C₂₀ alkynyl; C₁-C₂₀ haloalkyl; C₃-C₂₀ haloalkenyl; C₃-C₂₀ haloalkynyl; C₂-C₂₀ alkoxyalkyl; C₂-C₂₀ alkylthioalkyl; C₂-C₂₀ alkylsulfinylalkyl; C₂-C₂₀ alkylsulfonylalkyl; C₅-C₂₀ cycloalkylalkyl; C₄-C₂₀ alkenyloxyalkyl; C₄-C₂₀ alkynyloxyalkyl; C₄-C₂₀ (cycloalkyl) oxyalkyl; C₄-C₂₀ alkenylthioalkyl; C₄-C₂₀ alkynylthioalkyl; C₆-C₂₀ (cycloalkyl) thioalkyl; C₂-C₂₀ haloalkoxyalkyl; C₄-C₂₀ haloalkenyloxyalkyl; C₄-C₂₀ haloalkynyloxyalkyl; C₄-C₂₀ alkoxyalkenyl; C₄-C₂₀ alkoxyalkynyl; C₄-C₂₀ alkylthioalkenyl; C₄-C₂₀ alkylthioalkynyl; C₄-C₂₀ trialkylsilylalkyl; C₁-C₂₀ alkyl substituted with NR₃R₄, nitro, cyano, or phenyl optionally substituted with R₅, R₆, and R₇; C₁-C₂₀ alkoxy; C₁-C₂₀ haloalkoxy; C₁-C₂₀ alkylthio; C₁-C₂₀ haloalkylthio; NR₃R₄; or phenyl, benzyl, pyridyl, furanyl, thienyl, naphthyl, pyrimidinyl, benzofuranyl, benzothienyl, or quinolinyll each optionally substituted with R₅, R₆ or R₇;

R₃ is independently hydrogen; C₁-C₄ alkyl; or phenyl optionally substituted with at least one R₈;

R₄ is independently hydrogen; C₁-C₈ alkyl; or phenyl optionally substituted with at least one R₈;

R₅ is independently C₁-C₆ alkyl; C₁-C₆ alkoxy; C₁-C₆ haloalkyl; halogen; C₂-C₈ alkynyl; C₁-C₆ thioalkyl; phenyl or phenoxy each optionally substituted with at least one R₈; cyano; nitro; C₁-C₆ haloalkoxy; C₁-C₆ haloalkylthio; C₂-C₆ alkenyl; C₂-C₆ haloalkenyl; acetyl; CO₂CH₃; or N(C₁-C₂ alkyl)₂;

R₆ is independently methyl; ethyl; methoxy; methylthio; halogen; or trifluoromethyl;

R₇ is independently halogen; and

R₈ is independently halogen; C₁-C₄ alkyl; C₁-C₄ alkoxy; C₁-C₄ haloalkyl; nitro; or cyano;

wherein:

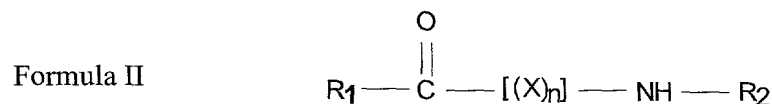
the amino acid in the first position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, lysine, methionine, serine, threonine and tryptophan;

the amino acid in the second position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, histidine, cysteine, threonine, tyrosine, and tryptophan; and

the amino acids in positions three through six, based on numbered amino acids from N-terminus to C-terminus, are any amino acid;

wherein the first two amino acids of said hexapeptides are other than arginine arginine, tryptophan-tryptophan, tryptophan-cysteine, tryptophan-lysine, arginine-tryptophan, or threonine-arginine.

32. An antimicrobial composition comprising a plurality of peptides, wherein said peptides each are represented by Formula II:



wherein:

X represents any amino acid except glutamate or aspartate;

n = 1-10;

R₁ is C₁-C₂₀ alkyl; C₃-C₆ cycloalkyl; C₄-C₂₀ alkenyl; C₄-C₂₀ alkynyl; C₁-C₂₀ haloalkyl; C₃-C₂₀ haloalkenyl; C₃-C₂₀ haloalkynyl; C₂-C₂₀ alkoxyalkyl; C₂-C₂₀ alkylthioalkyl; C₂-C₂₀ alkylsulfanylalkyl; C₂-C₂₀ alkylsulfonylalkyl; C₅-C₂₀ cycloalkylalkyl; C₄-C₂₀ alkenyloxyalkyl; C₄-C₂₀ alkynyloxyalkyl; C₄-C₂₀ (cycloalkyl) oxyalkyl; C₄-C₂₀ alkenylthioalkyl; C₄-C₂₀ alkynylthioalkyl; C₆-C₂₀ (cycloalkyl) thioalkyl; C₂-C₂₀ haloalkoxyalkyl; C₄-C₂₀ haloalkenyloxyalkyl; C₄-C₂₀ haloalkynyloxyalkyl; C₄-C₂₀ alkoxyalkenyl; C₄-C₂₀ alkoxyalkynyl; C₄-C₂₀ alkylthioalkenyl; C₄-C₂₀ alkylthioalkynyl; C₄-C₂₀ trialkylsilylalkyl; C₁-

C₂₀ alkyl substituted with NR₃R₄, nitro, cyano, or phenyl optionally substituted with R₅, R₆, and R₇; C₁-C₂₀ alkoxy; C₁-C₂₀ haloalkoxy; C₁-C₂₀ alkylthio; C₁-C₂₀ haloalkylthio; NR₃R₄; or phenyl, benzyl, pyridyl, furanyl, thienyl, naphthyl, pyrimidinyl, benzofuranyl, benzothienyl, or quinolinyl each optionally substituted with R₅, R₆ or R₇;

R₂ is C₁-C₂₀ alkyl; C₃-C₆ cycloalkyl; C₄-C₂₀ alkenyl; C₄-C₂₀ alkynyl; C₁-C₂₀ haloalkyl; C₃-C₂₀ haloalkenyl; C₃-C₂₀ haloalkynyl; C₂-C₂₀ alkoxyalkyl; C₂-C₂₀ alkylthioalkyl; C₂-C₂₀ alkylsulfinylalkyl; C₂-C₂₀ alkylsulfonylalkyl; C₅-C₂₀ cycloalkylalkyl; C₄-C₂₀ alkenyloxyalkyl; C₄-C₂₀ alkynyloxyalkyl; C₄-C₂₀ (cycloalkyl) oxyalkyl; C₄-C₂₀ alkenylthioalkyl; C₄-C₂₀ alkynylthioalkyl; C₆-C₂₀ (cycloalkyl) thioalkyl; C₂-C₂₀ haloalkoxyalkyl; C₄-C₂₀ haloalkenyloxyalkyl; C₄-C₂₀ haloalkynyloxyalkyl; C₄-C₂₀ alkoxyalkenyl; C₄-C₂₀ alkoxyalkynyl; C₄-C₂₀ alkylthioalkenyl; C₄-C₂₀ alkylthioalkynyl; C₄-C₂₀ trialkylsilylalkyl; C₁-C₂₀ alkyl substituted with NR₃R₄, nitro, cyano, or phenyl optionally substituted with R₅, R₆, and R₇; C₁-C₂₀ alkoxy; C₁-C₂₀ haloalkoxy; C₁-C₂₀ alkylthio; C₁-C₂₀ haloalkylthio; NR₃R₄; or phenyl, benzyl, pyridyl, furanyl, thienyl, naphthyl, pyrimidinyl, benzofuranyl, benzothienyl, or quinolinyl each optionally substituted with R₅, R₆ or R₇;

R₃ is independently hydrogen; C₁-C₄ alkyl; or phenyl optionally substituted with at least one R₈;

R₄ is independently hydrogen; C₁-C₈ alkyl; or phenyl optionally substituted with at least one R₈;

R₅ is independently C₁-C₆ alkyl; C₁-C₆ alkoxy; C₁-C₆ haloalkyl; halogen; C₂-C₈ alkynyl; C₁-C₆ thioalkyl; phenyl or phenoxy each optionally substituted with at least one R₈; cyano; nitro; C₁-C₆ haloalkoxy; C₁-C₆ haloalkylthio; C₂-C₆ alkenyl; C₂-C₆ haloalkenyl; acetyl; CO₂CH₃; or N(C₁-C₂ alkyl)₂;

R₆ is independently methyl; ethyl; methoxy; methylthio; halogen; or trifluoromethyl;

R₇ is independently halogen; and

R₈ is independently halogen; C₁-C₄ alkyl; C₁-C₄ alkoxy; C₁-C₄ haloalkyl; nitro; or cyano;

wherein:

the amino acid in the first position, based on numbered amino acids from N-terminus to

C-terminus, is selected from the group consisting of arginine, lysine, methionine, serine, threonine and tryptophan;

the amino acid in the second position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, histidine, cysteine, threonine, tyrosine, and tryptophan; and

the amino acids in positions three through six, based on numbered amino acids from N-terminus to C-terminus, are any amino acid.

33. The antimicrobial composition of claim 31 further comprising a carrier selected from the group consisting of a pharmaceutically acceptable carrier, an industrially acceptable carrier, a household product, and a personal care composition.

34. The antimicrobial composition of claim 32 further comprising a carrier selected from the group consisting of a pharmaceutically acceptable carrier, an industrially acceptable carrier, a household product, and a personal care composition.